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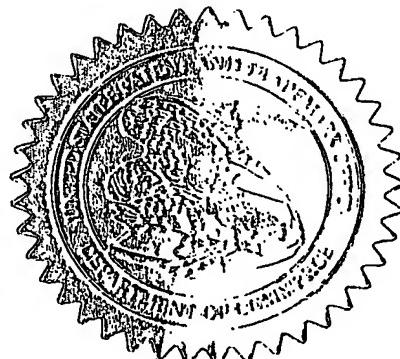
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PROVISIONAL APPLICATION COVER SHEET

Please request for filing a PROVISIONAL APPLICATION under 35 CFR 1.53 (c).

| | | | |
|---------------|--------|--------------------------------------|---|
| Docket Number | 60,349 | Type a plus sign (+) inside this box | + |
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TITLE OF THE INVENTION (280 characters max)

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Yes, the name of the U.S. Government agency and the Government contract number are:

Respectfully submitted,

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PROVISIONAL APPLICATION FILING ONLY

60349

**COMPOUNDS USEFUL AS INSECTICIDES
AND COMPOUNDS USEFUL AS ACARICIDES**

FIELD OF THE INVENTION

- 5 This invention provides compounds that are useful as insecticides and compounds that are useful as acaricides.

BACKGROUND OF THE INVENTION

- There is an acute need for new insecticides and acaricides. Insects and mites are developing resistance to the insecticides and acaricides in current use. At least 400 species of arthropods are resistant to one or more insecticides. The development of resistance to some of the older insecticides, such as DDT, the carbamates, and the organophosphates, is well known. But resistance has even developed to some of the newer pyrethroid insecticides and acaricides. Therefore, a need exists for new insecticides and acaricides, and particularly for compounds that have new or atypical modes of action.

DETAILED DESCRIPTION OF THE INVENTION

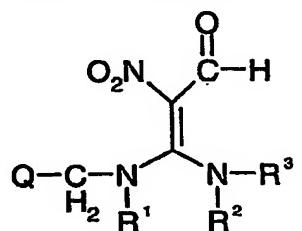


Figure One

- 20 In Figure One Q, R¹, R², and R³ have the following meanings. Q can be any five or six membered carbocyclic or heterocyclic ring, such as, phenyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl, pyrrolyl, pyrazolyl, imidazolyl, thienyl, furanyl, thiazolyl, isothiazolyl, oxazolyl, and isoxazolyl, and including reduced forms of the heterocyclic rings such as 25 tetrahydrofuranyl.

60349

R^1 , R^2 , and R^3 each independently can be:

- (a) a C₁₋₁₀, branched or unbranched, alkyl, alkoxy, alkenyl, alkynyl, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylcarbonyl, alkylcarbonothioyl, alkoxy carbonyl, alkylthiocarbonyl, alkoxy carbonothioyl, alkylthiocarbonothioyl, or HC(=NH)-;
- 5 (b) a C₃₋₁₀, cycloalkyl, or cycloalkenyl;
- (c) an aryl, heterocyclyl, aryloxy, heterocyclyloxy, arylthio, heterocyclithio, arylamino, or heterocyclylamino; or
- (d) a hydro, hydroxy, mercapto, amino, cyano, formyl, nitro, halo, 10 or aminocarbonyl.

Additionally, R^1 and R^2 can be joined together to form a ring, either directly with a bond between them, or indirectly through one or two linkage atoms, where such linkage atoms are either carbon, nitrogen, oxygen, or sulfur.

- 15 15 Each member of Q, each member of R^1 , R^2 , and R^3 , and any of the linkage atoms, which may have a hydrogen atom in a certain position, may instead of having such hydrogen atom, have a:

- (a) a C₁₋₁₀, branched or unbranched, alkyl, alkoxy, alkenyl, alkynyl, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylcarbonyl, alkylcarbonothioyl, alkoxy carbonyl, alkylthiocarbonyl, alkoxy carbonothioyl, alkylthiocarbonothioyl, HC(=NH)-, dialkylphosphonyl, or dialkylphosphatyl;
- 20 (b) a C₃₋₁₀, cycloalkyl, or cycloalkenyl;
- (c) an aryl, heterocyclyl, aryloxy, heterocyclyloxy, arylthio, heterocyclithio, arylamino, or heterocyclylamino; or
- 25 (d) a hydro, hydroxy, mercapto, amino, cyano, formyl, nitro, halo, or aminocarbonyl;

in such position, provided that these substituents are sterically compatible and the rules of chemical bonding and strain energy are satisfied.

60349

The term "aryl" means a monovalent radical derived by loss of hydrogen from an aromatic hydrocarbon. The term heteroaryl means a monovalent radical derived by loss of a hydrogen from a ring structure, where such ring structure contains one or more nitrogen, oxygen, or sulfur atoms. Examples of aryls and heteroaryls include, but are not limited to, phenyl, naphthyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl, pyrrolyl, pyrazolyl, imidazolyl, thienyl, furanyl, thiazolyl, isothiazolyl, oxazolyl, and isoxazolyl, and included are reduced forms of the heteroaryls such as tetrahydrofuranyl.

10 All salts and esters of these compounds are contemplated as part of this invention.

The compounds of the invention are useful for the control of pests such as, insects, mites, and aphids. Therefore, the present invention also is directed to a method for inhibiting an insect, mite, or aphid which 15 comprises applying to a locus of the insect or mite an insect- or mite-inhibiting amount of a compound of this invention. In particular, these compounds control insects in the order Homoptera, including the families Aphididae (aphids), Aleyrodidae (whiteflies), Delphacidae (planthoppers), and Cicadellidae (leafhoppers). They also control insects in the order 20 Coleoptera (beetles), including the family Chrysomelidae (leaf beetles). Also, they control lepidopterans such as tobacco bud worm and cabbage looper.

The compounds are useful for reducing populations of insects and mites and are useful in a method of inhibiting an insect or mite population 25 which comprises applying to a locus of the insect or mite an effective insect- or mite-inactivating amount of a compound of this invention.

The "locus" of insects or mites is a term used herein to refer to the environment in which the insects or mites live or where their eggs are present, including the air surrounding them, the food they eat, or objects

60349

or materials which they contact. For example, plant-ingesting insects or mites can be controlled by applying the active compound to plant parts that the insects or mites eat, particularly the foliage. Soil-inhabiting insects such as termites can be controlled by applying the active compound to the 5 soil that the insects move through. Insects such as fleas that infest animals can be controlled by applying the active compound to the animal that is infested.

It is contemplated that the compounds might also be useful to protect textiles, paper, stored grain, or seeds by applying an active compound to 10 such substance.

The term "inhibiting an insect or mite" refers to a decrease in the numbers of living insects or mites, or a decrease in the number of viable insect or mite eggs. The extent of reduction accomplished by a compound depends, of course, upon the application rate of the compound, the 15 particular compound used, and the target insect or mite species. At least an inactivating amount should be used.

The terms "insect-inactivating amount" and "mite-inactivating amount" are used to describe the amount, which is sufficient to cause a measurable reduction in the treated insect or mite population. Generally 20 an amount in the range from about 1 to about 1000 ppm by weight active compound is used. In another embodiment, the present invention is directed to a method for inhibiting a mite or aphid which comprises applying to a plant an effective mite- or aphid- inactivating amount of a compound of this invention.

25 The compounds of this invention are applied in the form of compositions which comprise a compound of this invention and a phytologically-acceptable inert carrier. The compositions are either concentrated formulations which are dispersed in water for application, or are dust or granular formulations which are applied without further

60349

treatment. The compositions are prepared according to procedures and formulae which are conventional in the agricultural chemical art, but which are novel and important because of the presence therein of the compounds of this invention.

- 5 The dispersions in which the compounds are applied are most often aqueous suspensions or emulsions prepared from concentrated formulations of the compounds. Such water-soluble, water-suspendable or emulsifiable formulations are either solids, usually known as wettable powders, or liquids usually known as emulsifiable concentrates or
10 aqueous suspensions. Wettable powders, which may be compacted to form water dispersible granules, comprise an intimate mixture of the active compound, an inert carrier, and surfactants. The concentration of the active compound is usually from about 10% to about 90% by weight. The inert carrier is usually chosen from among the attapulgite clays, the
15 montmorillonite clays, the diatomaceous earths, or the purified silicates.

Effective surfactants, comprising from about 0.5% to about 10% of the wettable powder, are found among the sulfonated lignins, the condensed naphthalenesulfonates, the naphthalenesulfonates, the alkylbenzenesulfonates, the alkyl sulfates, and nonionic surfactants such as ethylene
20 oxide adducts of alkyl phenols.

Emulsifiable concentrates of the compounds comprise a convenient concentration of a compound, such as from about 50 to about 500 grams per liter of liquid, equivalent to about 10% to about 50%, dissolved in an inert carrier which is either a water miscible solvent or a mixture of water-
25 immiscible organic solvent and emulsifiers. Useful organic solvents include aromatics, especially the xylenes, and the petroleum fractions, especially the high-boiling naphthalenic and olefinic portions of petroleum such as heavy aromatic naphtha. Other organic solvents may also be used, such as the terpenic solvents including rosin derivatives, aliphatic

60349

ketones such as cyclohexanone, and complex alcohols such as 2-ethoxyethanol. Suitable emulsifiers for emulsifiable concentrates are chosen from conventional nonionic surfactants, such as those discussed above.

- 5 Aqueous suspensions comprise suspensions of water-insoluble compounds of this invention, dispersed in an aqueous vehicle at a concentration in the range from about 5% to about 50% by weight. Suspensions are prepared by finely grinding the compound, and vigorously mixing it into a vehicle comprised of water and surfactants
10 chosen from the same types discussed above. Inert ingredients, such as inorganic salts and synthetic or natural gums, may also be added, to increase the density and viscosity of the aqueous vehicle. It is often most effective to grind and mix the compound at the same time by preparing the aqueous mixture, and homogenizing it in an implement such as a sand mill, ball mill, or piston-type homogenizer.
15

- The compounds may also be applied as granular compositions, which are particularly useful for applications to the soil. Granular compositions usually contain from about 0.5% to about 10% by weight of the compound, dispersed in an inert carrier which consists entirely or in large
20 part of clay or a similar inexpensive substance. Such compositions are usually prepared by dissolving the compound in a suitable solvent and applying it to a granular carrier which has been pre-formed to the appropriate particle size, in the range of from about 0.5 to 3 mm. Such compositions may also be formulated by making a dough or paste of the
25 carrier and compound and crushing and drying to obtain the desired granular particle size.

Dusts containing the compounds are prepared simply by intimately mixing the compound in powdered form with a suitable dusty agricultural

60349

carrier, such as kaolin clay, ground volcanic rock, and the like. Dusts can suitably contain from about 1% to about 10% of the compound.

The active compositions may contain adjuvant surfactants to enhance deposition, wetting and penetration of the compositions onto the target 5 crop and organism. These adjuvant surfactants may optionally be employed as a component of the formulation or as a tank mix. The amount of adjuvant surfactant will vary from 0.01 percent to 1.0 percent v/v based on a spray-volume of water, preferably 0.05 to 0.5 percent. Suitable adjuvant surfactants include ethoxylated nonyl phenols, 10 ethoxylated synthetic or natural alcohols, salts of the esters of sulphosuccinic acids, ethoxylated organosilicones, ethoxylated fatty amines, crop oil concentrates containing high molecular weight paraffinic oils and blends of surfactants with mineral and vegetable oils.

It is equally practical, when desirable for any reason, to apply the 15 compound in the form of a solution in an appropriate organic solvent, usually a bland petroleum oil, such as the spray oils, which are widely used in agricultural chemistry.

Insecticides and acaricides are generally applied in the form of a dispersion of the active ingredient in a liquid carrier. It is conventional to 20 refer to application rates in terms of the concentration of active ingredient in the carrier. The most widely used carrier is water.

The compounds of the invention can also be applied in the form of an aerosol composition. In such compositions the active compound is dissolved or dispersed in an inert carrier, which is a pressure-generating 25 propellant mixture. The aerosol composition is packaged in a container from which the mixture is dispensed through an atomizing valve. Propellant mixtures comprise either low-boiling halocarbons, which may be mixed with organic solvents, or aqueous suspensions pressurized with inert gases or gaseous hydrocarbons.

60349

The actual amount of compound to be applied to loci of insects, mites, and aphids is not critical and can readily be determined by those skilled in the art in view of the examples above. In general, concentrations of from 10 ppm to 5000 ppm by weight of compound are expected to provide 5 good control. With many of the compounds, concentrations of from 100 to 1500 ppm will suffice.

The locus to which a compound is applied can be any locus inhabited by an insect or arachnid, for example, vegetable crops, fruit and nut trees, grape vines, and ornamental plants.

10 Because of the unique ability of mite eggs to resist toxicant action, repeated applications may be desirable to control newly emerged larvae, as is true of other known acaricides.

In addition to being effective against mites, aphids, and insects when applied to foliage, compounds of this invention have systemic activity.

15 Accordingly, another aspect of the invention is a method of protecting a plant from insects which comprises treating plant seed prior to planting it, treating soil where plant seed is to be planted, or treating soil at the roots of a plant after it is planted, with an effective amount of a compound of this invention.

20 The action of the inventive compounds can be broadened by adding other, for example insecticidally, acaricidally, and/or nematocidally active, ingredients. For example, one or more of the following compounds can suitably be combined with the compounds of the invention:

25 (1) organophosphorus compounds such as acephate, azinphosmethyl, cadusafos, chlorethoxyfos, chlorpyrifos, coumaphos, dematon, demeton-S-methyl, diazinon, dichlorvos, dimethoate, EPN, ethoate, ethoprophos, etrimfos, fenamiphos, fenitrothion, fensulfothion, fenthion, fonofos, formothion, fosthiazate, heptenophos, malathion, methamidophos, methyl

60349

- parathion, mevinphos, monocrotophos, parathion, phorate, phosalone, phosmet, phosphamidon, phosphocarb, phoxim, profenofos, propaphos, propetamphos, prothiofos, pyrimiphos-methyl, pyrimiphos-ethyl, quinalphos, sulprofos; tebupirimphos, temephos, terbufos, tetrachlorvinphos, thiafenoxy, thiometon, triazophos, and trichlorphon;
- 5 (2) carbamates such as aldicarb, bendiocarb, benfuracarb, bensultap, BPMC, butoxycarbocim, carbaryl, carbofuran, carbosulfan, cloethocarb, ethiofencarb, fenobucarb, furathiocarb, methiocarb, isopropcarb, methomyl, oxamyl, pirimicarb, promecarb, propoxur, thiodicarb, and thifurox;
- 10 (3) pyrethroids such as acrinathrin, allethrin, beta-cyfluthrin, bifenthrin, bioresmethrin, cyfluthrin; cyhalothrin; lambda-cyhalothrin; gamma-cyhalothrin, cypermethrin; alpha-cypermethrin; zeta-cypermethrin; deltamethrin, esfenvalerate, fenvalerate, fenfluthrin, fenpropathrin, flucythrinate, flumethrin, fluvalinate, tau-fluvalinate, halfenprox, permethrin, protrifenbuta, resmethrin, silafluofen, tefluthrin, tetramethrin, tralomethrin, fish safe pyrethroids for example ethofenprox, natural pyrethrin, 15 tetramethrin, s-bioallethrin, fenfluthrin and prallethrin;
- 15 (4) acylureas, other types of insect growth regulators and insect hormone analogs such as buprofezin, chromfenozide, chlorfluazuron, diflubenzuron, fenoxy carb, flufenoxuron, halofenozide, hexaflumuron, hydroprene, leufenuron, 20 methoprene, methoxyfenozide, novaluron, pyriproxyfen, teflubenzuron and tebufenozide, N-[3,5-dichloro-2-fluoro-4-(1,1,2,3,3,3-hexafluoropropoxy)phenyl]-N'(2,6-difluorobenzoyl)urea;

60349

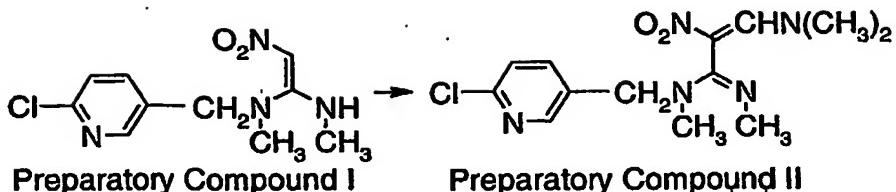
- (5) neonicotnioids and other nicotinics such as acetamiprid, AKD-1022, cartap, TI-435, clothianidin, MTI-446, dinotefuran, imidacloprid, nicotine, nitenpyram, thiamethoxam, thiacycloprid;
- 5 (6) macrolides such as avermectins, milbemycins, or spinosyns for example such as abamectin, ivermectin, milbemycin, emamectin benzoate and spinosad; and
- 10 (7) other insecticidal, acaricidal, mollscicial and nematocidal compounds or actives such as aldrin, amitraz, azadirachtin, azocyclotin, bifenazate, bromopropylate, chlordimeform, chlorgfenapyr, chlofentezine, chlorobenzilate, chlordane, cyhexatin, cyromazin, DDT, dicofol, dieldrin, DNOC, endosulfan, ethoxazole, fenazaquin, fenbutatin oxide, fenproximate, beta-fenpyroximate, fipronil, flubenzimine, hexythiazox, IKI-220, indoxacarb, lindane, methiocarb, metaldehyde, methoxychlor,
- 15 neem, petroleum and vegetable oils, pyridaben, pymetrozine, pyrimidifen, rotenone, S-1812, S-9539, spirodiclofen, sulfur, tebufenpyrad, tetradifon, triazamate, an insect-active extract from a plant; a preparation containing insect-active nematodes, a preparation obtainable from *Bacillus subtilis*, *Bacillus thuringiensis*, a nuclear polyhedrosis virus, or other like organism genetically modified or native, as well as synergists such as piperonyl butoxide, sesamax, safroxan and dodecyl imidazole, and phagostimulants such as cucurbitacin, sugars and Coax.
- 20

EXAMPLES

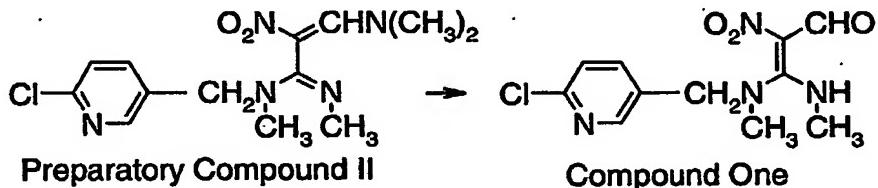
25 These examples are provided to further illustrate the invention. They are not meant to be construed as limiting the invention.

Example One: Preparation of Compound One

60349



- A solution of 6.0 g (23.4 mmol) of Preparatory Compound I (J. Pesticide Sci., 18, 31-40, 1983) in 35 mL of dimethylformamide dimethylacetal was heated at 85-90 °C for 3 hours and was then allowed 5 to cool to room temperature, which was about 22 °C. The volatiles were removed *in vacuo* and the resulting oil was triturated under ethyl ether and ethyl acetate with a small amount of methanol to give 2.2 g (30%) of Preparatory Compound II as a tan solid.



- 10 A solution of 130 mg (0.42 mmol) of Preparatory Compound II in 4 mL of concentrated ammonium hydroxide was stirred at room temperature, which was about 22°C, for 20 hours. The resulting mixture was concentrated to a residue which was chromatographed on silica gel (230-400 mesh) eluting with 95/5 dichloromethane/methanol to give 80 mg 15 (67%) of Compound One.

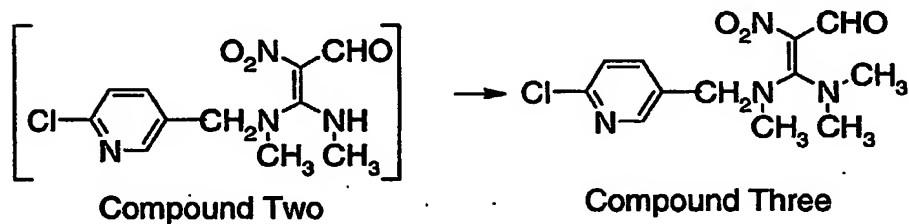
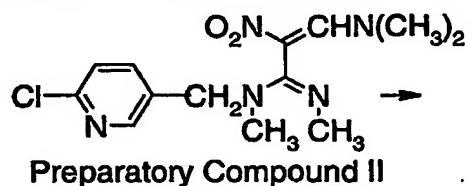
Example Two: Alternative Preparation of Compound One

- A solution of 200 mg (0.641 mmol) of Preparatory Compound II in 3 mL of methanol was treated with 0.32 mL (0.64 mmol) of 2.0 N sodium hydroxide. This mixture was then heated at reflux for 2.5 hours, followed 20 by cooling to about 22°C. The resulting portion was then concentrated to an oil, which was then treated with 0.64 mL (0.64 mmol) of 1.0 N hydrochloric acid. The resulting mixture was then extracted three times

60349

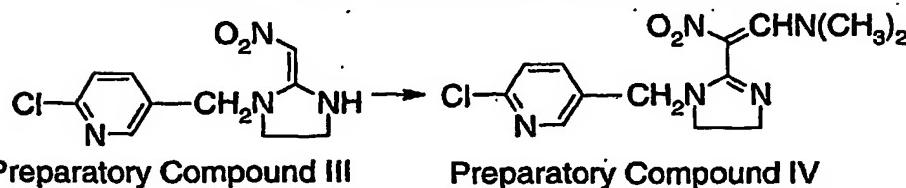
with dichloromethane, the combined extracts were then dried over sodium sulfate and were then concentrated to give 190 mg of a residue which was chromatographed on silica gel to give 130 mg (71%) of Compound One, mp 100-103 °C.

Example Three: Preparation of Compounds Two and Three



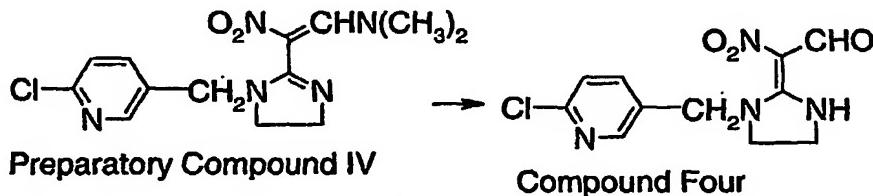
To a solution of 18.9 mg (0.822 mmol) of sodium metal in 3 mL of absolute ethanol was added in one portion 200 mg (0.641mmol) of Preparatory Compound II. The contents were stirred at room temperature, which was about 22 °C, for 3 hours. The solution was cooled in ice and was treated with 0.13 mL (296 mg, 2.09 mmol)) of methyl iodide in 1.5 mL of ethanol. After stirring overnight at room temperature, which was about 22 °C, the mixture was concentrated and the residue was partitioned between dichloromethane and water. Hydrochloric acid (1.0 N, 0.64 mL) was added followed by sodium bicarbonate to give a pH of 10-11. The mixture was extracted three times with dichloromethane, the combined extracts were dried over magnesium sulfate and were concentrated to give 160 mg of an oil which was chromatographed on silica gel using 95/5 dichloromethane/methanol as eluant to give 10 mg (5%) of Compound Three, mp 163-166 °C and 130 mg (71%) of Compound Two.

60349

Example Four: Preparation of Compound Four

- A solution of 255 mg (1.00 mmol) of Preparatory Compound III (EP 163855 A1) and 238 mg (2.00 mmol) of dimethylformamide
 5 dimethylacetal in 3 mL of dry toluene was heated at 100-110 °C for 6 hours and was then allowed to cool to room temperature, which was about 22°C. The toluene was removed *in vacuo* and the resulting solid was triturated under ethyl ether to afford 272 mg (88%) of Preparatory Compound IV as a yellow solid, mp 115-123 °C.

10



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- A solution of 161 mg (0.521 mmol) of Preparatory Compound IV and 0.286 mL (0.573 mmol) of 2.0 N sodium hydroxide in 3 mL of methanol was stirred at room temperature, which was about 22°C, for 7 hours and was then cooled in ice. The resulting solution was then treated with 0.572 mL (0.572 mmol) of 1.0 N hydrochloric acid. The precipitate was collected to afford 90 mg (61%) of Compound Four, mp 196-197 °C (dec).

Example Five: Testing with Cotton Aphid (*Aphis gossypii*)

20

- Alcala cotton plants were grown from seed in 7-cm pots under greenhouse conditions until they reached 35 to 40 cm in height; approximately four weeks old with 5 to 6 true leaves. The plants were then stripped of all foliage except for the two uppermost true leaves. Two days prior to application, heavily infested leaf sections from a cotton aphid colony were cut and placed on each untreated leaf surface. Over the two

60349

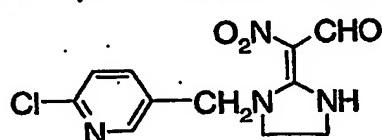
days, all stages of the aphids abandon the excised host material and migrate to the succulent growth, predominantly the lower surface of the leaves. Plants were checked prior to application for even infestation levels. Treatments consisted of 4 replicates (plants) each with 2 leaves per plant.

Formulation of technical compound was in a solvent solution. For the top rate of each compound, 20 milligrams of technical material was dissolved in 2 milliliters (ml) of a 9:1 mixture of acetone:ethanol solvent. Once dissolved, an additional 18-ml of the solvent mix was added to yield a 1 mg/ml spray solution. Additional rates were then prepared by serial dilution.

Application was made by an automated tracksprayer (RC-Insecticide #1, Serial MS-9L2-002, MANDEL Scientific Company Ltd.) equipped with a single TX6 SS hollow cone nozzle to simulate an over-the-top application by a boom sprayer. Spray pressure was set at 50 PSI, and the nozzle head was set approximately 40 cm above the leaf surface (depending on the leaf position it ranged from 35-45 cm). Track velocity and all other settings were calibrated to deliver approximately 200L/ha. After application and drying, the plants were transferred and kept in controlled environment until assessment.

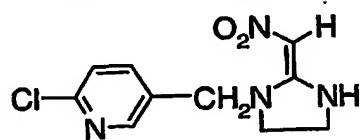
Assessment of compound efficacy is evaluated by counting all living non-winged stage aphids infesting each replicate using a dissecting microscope.

The compounds tested in this manner were as follows.



25

Compound Four



Comparative Compound A

60349

The results indicate that the LC90 in grams per hectare for Compound Four was less than 0.78 grams per hectare, whereas, the L90 for Comparative Compound A was 9.1 grams per hectare. This means nearly 11.7 times more Comparative Compound A was needed to reach the 5 LC90 level than Compound Four.

Example Six: Testing with Sweetpotato Whitefly, *Bemisia tabaci*

Greenhouse-grown cotton plants (ca. five weeks old, grown in 7 x 7 cm pots containing ca. 231 cm³ of potting soil mixture) were prepared for whitefly infestation by removing all foliage except the first or second pair 10 of fully expanded true leaves. These plants are then moved into the whitefly colony room of the insectary (temperature maintained at 27° C), where they are surrounded by cotton or velvetleaf plants infested with all life stages of *B. tabaci*. Initial movement of *B. tabaci* adults onto the prepared cotton plants is encouraged by gently shaking the foliage of the 15 surrounding infested cotton and velvetleaf foliage using a 1.5-meter long willow branch. The prepared plants are left in place within the whitefly colony, where they are exposed to adult female *B. tabaci* oviposition for ca. 48 hours.

At the end of the oviposition exposure period, adult whiteflies are 20 dislodged from the prepared plants by passing compressed air just above the leaves while the leaves are gently disturbed by hand for ca. 15 seconds. The prepared plants, which are now infested with eggs of *B. tabaci*, are removed from the insectary and any remaining adults are removed from the foliage with a hand-held vacuum fitted with a cage to 25 trap insects. Infested plants are then selected at random and arranged into treatment groups of two or three plants each. Leaves of plants to receive different treatments are separated by a distance of at least 5 cm.

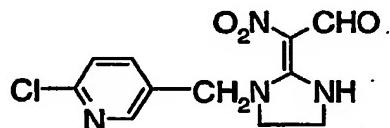
Compounds are applied within 24 hours of removing cotton plants from exposure to oviposition within the *B. tabaci* colony. Immediately after

60349

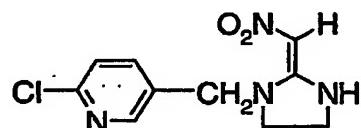
application, plants are moved to a holding room with a temperature of 30°C and relative humidity of ca. 60%. Compound efficacy was observed by an assessment at 13 days after application. At this evaluation time, whitefly nymphs that have successfully developed to the third and fourth nymphal stadia can be easily seen on the underside of each cotton leaf with the aid of an illuminated magnifying lens (equipped with a circular fluorescent tube).

Track sprayer application is designed to simulate field application of compounds. As in field applications, the equipment is first calibrated to deliver the desired spray volume, in liters/ha. Next, volumes of spray solution containing known amounts of test compound are prepared, so that when applied at the known liters/ha spray volume, the desired g [All]/ha rate will be delivered.

The compounds tested in this manner were as follows.



15 Compound Four



Comparative Compound A

The results indicate that the LC90 in grams per hectare for Compound Four was less than 2.52 grams per hectare, whereas, the L90 for Comparative Compound A was 6.66 grams per hectare. This means nearly 2.7 times more Comparative Compound A was needed to reach the LC90 level than Compound Four.

60349

WE CLAIM

1. A compound according to

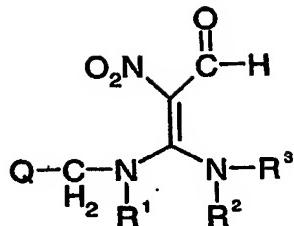


Figure One

5 wherein

Q can be any five or six membered carbocyclic or heterocyclic ring,
 R¹, R², and R³ each independently can be

- (a) a C₁₋₁₀, branched or unbranched, alkyl, alkoxy, alkenyl, alkynyl, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylcarbonyl, alkylcarbonothioly, alkoxy carbonyl, alkylthiocarbonyl, alkoxy carbonothioly, alkylthiocarbonothioly, HC(=NH)-,
- (b) a C₃₋₁₀, cycloalkyl, or cycloalkenyl,
- (c) an aryl, heterocycl, aryloxy, heterocyclxy, arylthio, heterocyclthio, arylamino, or heterocyclamino, or
- (d) a hydro, hydroxy, mercapto, amino, cyano, formyl, nitro, halo, or aminocarbonyl,

10 R¹ and R² can be joined together to form a ring, either directly with a bond between them, or indirectly through one or two linkage atoms, where such linkage atoms are either carbon, nitrogen, oxygen, or sulfur, and

20 wherein

each member of Q, each member of R¹, R², and R³, and any of the linkage atoms, which may have a hydrogen atom in a certain position, may instead of having such hydrogen atom, have a

- 25 (a) a C₁₋₁₀, branched or unbranched, alkyl, alkoxy, alkenyl,

60349

alkynyl, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylcarbonyl, alkylcarbonothioyl, alkoxy carbonyl, alkylthiocarbonyl, alkoxy carbonothioyl, alkylthiocarbonothioyl, HC(=NH)-, dialkylphosphonyl, or dialkylphosphatyl,

- 5 (b) a C₃₋₁₀, cycloalkyl, or cycloalkenyl,
 (c) an aryl, heterocyclyl, aryloxy, heterocyclyoxy, arylthio, heterocyclithio, arylamino, or heterocyclamino, or
 (d) a hydro, hydroxy, mercapto, amino, cyano, formyl, nitro, halo, or aminocarbonyl,
- 10 in such position.

2. A composition comprising a compound according to claim 1 and at least one other active compound where such active compound is at least insecticidally, acaricidally, or nematocidally active.

- 15 3. A process of applying a compound according to claim 1, or a composition according to claim 2, to a locus in an amount effective to control pests.
- 20 4. A process of applying a compound according to claim 1, or a composition according to claim 2, to a locus in an amount effective to control insects or mites.

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ABSTRACT

Compounds useful to control pests are provided.

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